

INVITATION / TRAINING

# GOOD CLINICAL PRACTICE

## Refresher Training for investigators and their staff

In the framework of the Luxclin conference, CIEC organizes an interactive and certified training in “Good Clinical Practice”, focusing on updated ICH GCP guidelines (E6 R2 Update).

### AGENDA

<b>13:30-13:40</b>	Welcome and Introduction
<b>13:40-14:00</b>	What is new in the ICH-GCP Guidelines with Integrated Addendum E6 (R2)? <i>(Ingrid Klingmann)</i>
<b>14:00-14:45</b>	Sponsor responsibility: Focus on quality management <i>(Ingrid Klingmann)</i>
<b>14:45-15:15</b>	Break
<b>15:15-16:00</b>	Practical aspects of a risk-based monitoring approach <i>(Ingrid Klingmann)</i>
<b>16:00-17:00</b>	Site management with validated computerized systems in Europe and US (CRF part 11) <i>(Lisbeth Tofte)</i>
<b>17:00-17:30</b>	Final test
<b>17.30-17:45</b>	Review and discussion of correct answers <i>(Ingrid Klingmann and Lisbeth Tofte)</i>
<b>17:45</b>	Closing remarks



Speakers: **Ingrid Klingmann**, MD, PhD  
Chairman, European Forum for Good Clinical Practice (EFGCP)

**Lisbeth Tofte**, MD, Drug Safety Consult,  
Copenhagen, member of EFGCP

**SAVE THE DATE**  
**31st May 2017**  
**01:30 pm - 05:45 pm**

**VENUE**  
**Centre Hospitalier  
Luxembourg (CHL)**

Amphithéâtre  
4, rue Barblé  
L-1210 Luxembourg

**INFORMATION & REGISTRATION**  
Registration :

[gcptraining.lih.lu](http://gcptraining.lih.lu)

As the number of seats is limited, we will accept registrations on a first come first served basis.

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**IN COLLABORATION WITH**



**Luxclin conference**

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